



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) **EP 1 081 108 A1**

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
07.03.2001 Bulletin 2001/10

(51) Int. Cl.⁷: **C03C 17/23, C03C 17/245,
C03C 17/25**

(21) Application number: **00118871.3**

(22) Date of filing: **31.08.2000**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: **02.09.1999 JP 24815599
24.12.1999 JP 36712199**

(71) Applicant:
**Central Glass Company, Limited
Ube-shi, Yamaguchi 755-0001 (JP)**

(72) Inventors:
• **Yamamoto, Hideki,
c/o Glass Research Center of
Matsusaka-shi, Mie 515-0001 (JP)**

- **Yamazaki, Seiji,
c/o Glass Research Center of
Matsusaka-shi, Mie 515-0001 (JP)**
- **Nishida, Yoshihiro,
c/o Glass Research Center of
Matsusaka-shi, Mie 515-0001 (JP)**
- **Honjo, Hiroshi,
c/o Glass Research Center of
Matsusaka-shi, Mie 515-0001 (JP)**

(74) Representative:
**Manitz, Finsterwald & Partner
Postfach 22 16 11
80506 München (DE)**

(54) **Article with photocatalytic film**

(57) The invention relates to an article having a substrate and photocatalytic film covering the substrate. This film contains (1) a film-forming component including ZrO_2 and SiO_2 and (2) TiO_2 crystals dispersed in the film-forming component. These ZrO_2 , SiO_2 and TiO_2 crystals respectively are in amounts of 25-60 wt%, 15-50 wt% and 25-45 wt%, based on the weight of the film. The article is improved in photocatalytic activity, abrasion resistance and chemical resistance by the provision of the film.

EP 1 081 108 A1

Description**BACKGROUND OF THE INVENTION**

[0001] The present invention relates to an article with a photocatalytic film, such as an automotive or architectural window pane. The article is provided with anti-fogging function, stain resistance, hydrophilicity and good visibility in rainy weather.

[0002] Japanese Patent Laid-open Publication JP-A-9-328336 discloses a composition for forming a photocatalytic film. This composition comprises (1) TiO_2 grains having an average grain diameter of less than 100 nm, (2) Zr-containing compound, and (3) Si-containing compound. The composition is such that the weight ratio of the Zr-containing compound to the TiO_2 grains is from 0.02 to 0.5 in terms of oxide and that the weight ratio of the Si-containing compound to the TiO_2 grains is from 0.2 to 2.5 in terms of oxide.

[0003] JP-A-10-216528 discloses a photocatalytic article comprising a substrate and a catalytic film formed thereon. This catalytic film is a combination of photocatalytic grains and a binder containing 60-90 wt% of silica and 10-40 wt% of zirconia.

[0004] JP-A-9-227159 discloses a vehicular front or rear window glass having a substantially transparent layer bonded to a substrate. This layer contains a photocatalytic semiconductor material.

[0005] JP-A-10-156999 discloses a photocatalytic layer made of a mixture of silica and titania. JP-A-9-271731 discloses a photocatalytic member having a top surface comprising titanium oxide and silicon oxide as major components. JP-A-9-56788 discloses an article for bathroom, having a photocatalytic layer made of a mixture of silica and titania.

[0006] JP-A-11-35342 discloses a multi-functional glass having a soda-lime glass substrate, a first layer formed on the substrate, and a second layer formed on the first layer. The first layer is a SiO_2 film containing at least one member selected from Ti and metal elements having electronegativities close to that of Ti. The second layer is a TiO_2 film or a SiO_2 film in which TiO_2 fine particles are dispersed.

SUMMARY OF THE INVENTION

[0007] It is an object of the present invention to provide an article with a photocatalytic film, which is improved in photocatalytic activity, abrasion resistance and chemical resistance.

[0008] It is another object of the present invention to provide a process for producing such article.

[0009] According to the present invention, there is provided an article comprising a substrate; and a photocatalytic film covering said substrate. This film comprises (1) a film-forming component comprising ZrO_2 and SiO_2 and (2) TiO_2 crystals dispersed in said film-forming component, said ZrO_2 , said SiO_2 and said TiO_2 crystals respectively being in amounts of 25-60 wt%, 15-50 wt% and 25-45 wt%, based on a weight of said film.

[0010] According to the present invention, there is provided a first process for producing the article. The first process comprises (a) providing a coating liquid comprising a first raw material of said ZrO_2 , a second raw material of said SiO_2 , and a third raw material of said TiO_2 crystals; (b) applying said coating liquid to a surface of said substrate, thereby forming thereon a precursory film; and (c) subjecting said precursory film to a heat treatment at a temperature of 300-700°C, thereby turning said precursory film into said photocatalytic film.

[0011] According to the present invention, there is provided a second process for producing the article. The second process comprises (a) providing a coating liquid comprising a first raw material of said ZrO_2 , a second raw material of said SiO_2 , and, a third raw material of said TiO_2 crystals; (b) applying said coating liquid to a surface of said substrate, thereby forming thereon a precursory film; and (c) subjecting said precursory film to a first baking at a temperature of 560-700°C, thereby turning said precursory film into said photocatalytic film.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0012] We unexpectedly found that an article according to the invention is improved in photocatalytic activity (e.g., stain resistance and hydrophilicity), abrasion resistance and chemical resistance (e.g., acid resistance and alkali resistance) by specifying respective amounts of ZrO_2 , SiO_2 and TiO_2 crystals as above, particularly by increasing the amount of ZrO_2 relative to that of TiO_2 crystals as above. For example, the obtained article becomes satisfactory in an alkali resistance test for Category A of Japanese Industrial Standard (JIS) R 3221 and thus can practically be used as an exterior window pane such as automotive or architectural window pane.

[0013] As stated above, the respective amounts of ZrO_2 , SiO_2 and TiO_2 crystals are specified. Particularly, TiO_2 crystal fine grains, for example, having an average grain size of not greater than 30 nm, are preferably used as at least part of a raw material of TiO_2 crystals of the photocatalytic film. Due to these features, the photocatalytic film does not have cracks even if the baking of the precursory film is conducted at a high temperature at which a glass plate (substrate) can be bent into a curved glass plate. Generally speaking, such high temperature tends to make a photocatalytic

film inferior in photocatalytic activity. According to the invention, however, the photocatalytic film does not become inferior in photocatalytic activity by such high temperature, due to the above features. Therefore, it is not necessary to provide an alkali barrier layer between the glass plate and the photocatalytic film for preventing the movement of alkali components of the glass plate into the photocatalytic film.

- 5 [0014] The total amount of ZrO_2 , SiO_2 and the TiO_2 crystals of the photocatalytic film is preferably at least 90 wt%, based on the weight of the film, in order to have photocatalytic activity and durability such as abrasion resistance, acid resistance and alkali resistance. Besides these essential components, the film may optionally comprise not greater than 10 wt% of at least one component such as amorphous TiO_2 , colloidal silica, other oxides (e.g., Al_2O_3 , B_2O_3 and SnO_2), ions (Cr ion and V ion) and carbon. The TiO_2 crystals of the film may have an average grain size of not greater than 30 nm or not greater than 20 nm.

[0015] In case that the article is used as an automotive window pane, it is preferable to dispose the photocatalytic or hydrophilic film on an outer side of automobile.

- 10 [0016] In case that a glass plate is used as the substrate in the second process, the glass plate may be bent into a curved glass plate after the first baking or when the precursory film and the glass substrate are subjected to the first baking.

[0017] In order to improve the film in transparency and durability, the third raw material of the first process is preferably TiO_2 grains having an average grain size of not greater than 20 nm, and that of the second process is preferably those grains having an average grain size of not greater than 30 nm.

- 20 [0018] The first raw material of the first and second processes is preferably zirconium chloride or zirconium nitrate in order to improve the photocatalytic film in durability, particularly abrasion resistance, acid resistance and alkali resistance. Examples of the zirconium chloride include zirconium chloride, zirconium oxychloride octahydrate, and a chlorine-containing zirconium alkoxide represented by the general formula $\text{Zr}(\text{OC}_m\text{H}_{2m+1})_x\text{Cl}_y$, where m, x and y are integers and $x+y=4$. Examples of the zirconium nitrate include zirconium oxynitrate dihydrate. The ZrO_2 content of the film is not lower than 25 wt% in order to have the film provided with alkali resistance and photocatalytic activity. If it is higher than 60 wt%, the film becomes inferior in abrasion resistance and/or photocatalytic activity. The ZrO_2 content of the film is preferably from 25 to 45 wt%. As ZrO_2 is added to the film, the film is improved in alkali resistance. Furthermore, the film is remarkably improved in photocatalytic activity if the ZrO_2 content of the film is not lower than 25 wt%.

- 25 [0019] The second raw material of the first and second processes is not particularly limited, so long as it can produce SiO_2 after the heat treatment or the first baking. Examples of the second raw material include alkoxysilanes such as tetramethoxysilane, tetraethoxysilane, methyltrimethoxysilane and methyltriethoxysilane, hydrolysates of these, polycondensates of these, those stabilized with stabilizers such as acetylacetone, and commercial products for producing SiO_2 such as COLCOAT P of COLCOAT Co., MSH2 of Mitsubishi Chemical Corp. and CSG-DI-0600 of Chisso Co. The SiO_2 content of the film is not lower than 15 wt% in order to have the film provided with abrasion resistance. If it is higher than 50 wt%, the film becomes inferior in photocatalytic activity and/or alkali resistance. The SiO_2 content of the film is preferably from 25 to 45 wt%.

- 30 [0020] The third raw material of the first and second processes is not particularly limited, so long as it can produce TiO_2 crystals after the heat treatment or the first baking. It is possible to use commercial powdery TiO_2 fine grains for photocatalyst, such as ST-01 and ST-21 of Ishihara Techno Co., SSP-25 and SSP-20 of SAKAI CHEMICAL INDUSTRY CO., LTD., PC-101 of Titan Kogyo K.K., SUPER TITANIA F-6 and SUPER TITANIA F-5 of Showa Titanium Co., and DN-22A of FURUKAWA CO., LTD. It is further possible to use liquid-type commercial products of the third raw material, such as STS-01 and STS-02 of Ishihara Techno Co. and A-6 and M-6 of TAKI CHEMICAL CO., LTD. It is still further possible to use a mixture of TiO_2 fine grains for photocatalyst and a raw material of silica, such as ST-K01 and ST-K03 (trade names) of Ishihara Techno Co. The TiO_2 crystal content of the film is not lower than 25 wt% in order to have a photocatalytic capability to achieve hydrophilicity and stain resistance. It is not greater than 45 wt% from the viewpoint of alkali resistance and abrasion resistance. Powdery TiO_2 fine grains can easily be dispersed in a coating liquid by a common mixing operation for dispersing a powder in a liquid, for example, using a ball mill. Upon this, it is optional to disperse the first raw material of ZrO_2 and the second raw material of SiO_2 in the coating liquid.

[0021] The manner of applying the coating liquid to the surface of a substrate is not particularly limited. It can be conducted by a common method such as dip coating, spin coating, roller coating, spraying, screen printing or the like.

- 40 [0022] In the first process, the precursory film is subjected to a heat treatment at a temperature of 300-700°C, thereby turning the precursory film into the photocatalytic film. Prior to this heat treatment, it is optional to conduct a drying at a temperature not higher than 200°C or a preliminary baking at a temperature lower than that of the heat treatment. It is preferable to regulate the temperature of the heat treatment in a manner that a period of time, during which the heat treatment temperature lower than a maximum temperature of the heat treatment by 100°C or less, is at least 1 minute, preferably at least 3 minutes. This maximum temperature can be arbitrarily set within a range of 300-700°C. The upper limit of the heat treatment time is not particularly limited. For example, a heat treatment time of 24 hr does not cause particular problems. The above-mentioned period of time is preferably about 2 hr from the viewpoint of productivity.

[0023] In the second process, the precursory film is subjected to a first baking at a temperature of 560-700°C, preferably 600-700°C, thereby turning the precursory film into the photocatalytic film. During the first baking, the substrate may be bent if it is necessary to produce a curved substrate. The film is provided with durability by the first baking. It is optional to conduct a second baking (preliminary baking) at a temperature of 300-620°C, preferably 450-620°C, prior to the first baking (main baking). A film becomes more compact and improved particularly in abrasion resistance by conducting these two bakings. In case that an automotive curved window pane is produced as the article of the invention, it is preferable to regulate the temperature of the first baking in a manner that a period of time, during which the first baking temperature lower than a maximum temperature of the first baking by 100°C or less, is at least 1 minute, preferably at least 2 minutes. This maximum temperature can be arbitrarily set within a range of 560-700°C. The upper limit of the first baking time is not particularly limited. For example, the above-mentioned period of time is preferably about 3 minutes in case that a glass substrate is tempered by the first baking and preferably about 10 minutes in case that a glass substrate is bent during the first baking for producing a laminated glass. The TiO₂ crystals in the form of fine grains are dispersed in the film-forming component of the film, and the TiO₂ crystal content of the film is limited to a range of 25-45 wt%. With this, the film does not have cracks even if a glass substrate is bent during the first baking.

[0024] The film thickness is preferably from 30 to 500 nm in order to have photocatalytic activity and a high durability. It is more preferably from 50 to 300 nm in order to have a good transparency and a still high durability by the formation of a photocatalytic film one time. The material of the substrate is not particularly limited, as long as it does not deteriorate by heating. It may be selected from glass, ceramics and metals (e.g., aluminum and stainless steel). In particular, glass is preferable because of its transparency and heat resistance.

[0025] The article according to the invention is improved in durability, hydrophilicity, anti-fogging and stain resistance by the provision of the photocatalytic film. Thus, it can be used in various severe environments for building, vehicular, aircraft and industrial window panes, exterior tiles, exterior panels, showcases, and various mirrors such as vehicular door mirror and bathroom mirror. In particular, the article is preferably used for vehicular front, rear and side window panes. The article is improved in visibility under a rainy weather by the provision of hydrophilicity.

[0026] As mentioned above, the substrate can be a glass plate, for example, of a soda-lime-silicate glass. The glass plate is not limited to particular types. For examples, it may be a clear or colored glass plate having a color of blue, gray, bronze, green or the like, a glass reinforced with net or the like, a curved, semi-tempered or tempered glass, a double layer glass (sealed double-glazed unit) optionally having a metal film such as Low-E film, a double layer glass having a gel between two glass plates, a laminated glass, a glass plate pierced or formed on its uncoated side with a metal, oxide or resin film by deposition, sputtering, printing or the like, or a glass plate having an uncoated side subjected to etching, sand blasting or the like.

[0027] When the TiO₂ crystals of the photocatalytic film are irradiated with ultraviolet rays contained in the sunlight or a fluorescent light, organic contaminants on the film are oxidatively decomposed by the photocatalytic activity of the TiO₂ crystals. With this, the surface of the film becomes clean, and the surface of the TiO₂ crystals becomes hydrophilic. A film not according to the invention made of only TiO₂ crystals also becomes hydrophilic by its irradiation with ultraviolet rays. This film, however, becomes hydrophobic in a relatively short time due to the original hydrophobicity of TiO₂, if the irradiation is not repeated. In contrast, according to the invention, SiO₂ is added to the film. With this, the film is improved in hydrophilicity. Furthermore, a larger amount of water is adsorbed to the film, thereby improving the photocatalytic capability of the film. Still furthermore, the film is improved in durability such as abrasion resistance. According to the invention, ZrO₂ is added to the film. With this, the film is further improved in durability, particularly abrasion resistance and alkali resistance, and in photocatalytic capability. Thus, according to the invention, the photocatalytic film is very much improved in photocatalytic capability and durability, particularly abrasion resistance and alkali resistance. The surface of the photocatalytic film is kept hydrophilic. The film is capable of decomposing organic contaminants (e.g. organic components of exhaust gas and dust) by the photocatalytic activity. Furthermore, stain on the film runs down easily by rain water, since the film surface is kept hydrophilic. Furthermore, even if a glass plate used as the substrate is bent during the baking, the film does not have cracks and have a high quality and a high durability.

[0028] The following nonlimitative examples are illustrative of the present invention.

EXAMPLE 1-1

[0029] A coating liquid was prepared by dispersing, in a solvent that is a mixture of ethanol and 1-methoxy-2-propanol each being made of Kishida Chemical Co., (1) ZrOCl₂ of Kishida Chemical Co. as the raw material of ZrO₂, (2) CSG-DI-0600 (trade name) of Chisso Co., containing 6% of a siloxane oligomer in a mixture of ethanol and isopropanol, as the raw material of SiO₂, and (3) ST-K01 (trade name) of Ishihara Techno Co., in which the weight ratio of TiO₂ to SiO₂ is 8:2, in amounts such that the film contained 30 wt% of ZrO₂, 30 wt% of SiO₂ and 40 wt% of TiO₂. Separately, a float glass plate of soda-lime-silicate glass having widths of 100 mm and a thickness of 3mm was washed sufficiently with water, detergent and ceria. This glass plate was used as the substrate. The coating liquid was applied to the surface of the glass plate by spin coating. The coated glass plate was dried for 20 minutes in a DK43-type air-circulating

constant-temperature thermostat (made by Yamato Kagaku Co.) of 150°C, followed by a heat treatment for 10 minutes in a FP41-type muffle furnace (made by Yamato Kagaku Co.) of 650°C, thereby obtaining an article with a photocatalytic film having a thickness of 120 nm and containing 30 wt% of ZrO₂, 30 wt% of SiO₂ and 40 wt% of TiO₂, based on the weight of the film.

5 [0030] The obtained article (test sample) was subjected to the following abrasion resistance test, acid resistance test, alkali resistance test, photocatalytic activity test, and hydrophilicity test. The results of these tests are shown in Table 1. Of these, the evaluations of the abrasion resistance test, the acid resistance test and the alkali resistance test were conducted based on Category A of JIS R 3221.

10 [0031] The abrasion resistance test was conducted in accordance with JIS R 3221. The haze value was measured before and after an abrasion test. In this test, the test sample was rotated certain cycles relative to the Taber abraser having an abrasive wheel CS-10F, while the abrasive wheel is in abutment with the test sample under a load of 500 gf. In Table 1, "good" means that the haze value before the test H_0 , the haze value after the 100 cycles H_{100} , and the haze value after the 200 cycles H_{200} are in a relation of $H_0 \leq H_{100} \leq H_{200}$ and that ΔH is 4% or less where $\Delta H = H_{200} - H_0$. In Table 1, "not good" means that H_{100} is greater than H_{200} or ΔH is greater than 4%.

15 [0032] The acid resistance test was conducted in accordance with JIS R 3221. In this test, the test sample was immersed for 24 hr in 1N hydrochloric acid maintained at 23°C \pm 2°C. After that, the coated side of the test sample was wiped with a flannel cloth while a flowing water was applied to the test sample, followed by drying. Then, the external appearance of the test sample was observed with the naked eyes. In Table 1, "good" means that the film did not change substantially in its external appearance, and "not good" means that the film had a substantial coloration or scratches or
20 that the film exfoliated.

[0033] The alkali resistance test was conducted in the same manner as that of the acid resistance test, except that 1N hydrochloric acid was replaced with 1N sodium hydroxide aqueous solution.

[0034] In the photocatalytic activity test, the photocatalytic activity of the test sample was evaluated by the degree of decomposition of stearic acid on the film due to the ultraviolet irradiation. In this test, the test sample was immersed
25 in a solution containing 3wt% of stearic acid dissolved in ethanol, and then withdrawn from the solution at a rate of 8 mm/sec. After that, the test sample was irradiated for 1 hr with ultraviolet rays from a black light, FL15BLB (trade name) of Toshiba Denki Co. in a manner to have a ultraviolet strength of 4 mW/cm² (365 nm) at the coated surface of the test sample. The amount of stearic acid on the film was determined by measuring a peak strength (absorbance A) appearing within a range of 2910-2920 cm⁻¹ and derived from the C-H stretching vibration of stearic acid, using a FT-IR spec-
30 trophotometer of Perkin-Elmer Co., Ltd., before and after the application of stearic acid and after the ultraviolet irradiation for 1 hr. Then, a peak strength change as the degree of stearic acid decomposition was determined by the expression $\{(A_0 - A_b) - (A_1 - A_b)\} \times 1000$ where A_b is the peak strength before the application of stearic acid, A_0 is the peak strength after the application of stearic acid but before the ultraviolet irradiation, and A_1 is the peak strength after the ultraviolet irradiation for 1 hr. The results of the peak strength change are shown in Table 1. A peak strength change
35 of 5 or greater was judged to be satisfactory, and that of less than 5 unsatisfactory.

[0035] The hydrophilicity test was conducted by allowing the test sample to stand still for 7 days in a laboratory of an environment having an ultraviolet strength not greater than 1 μ W/cm² (365 nm). Then, a water drop was placed on the coated side of the test sample, and its contact angle was measured. In Table 1, "good" means that the contact angle was not greater than 20 degrees, and "not good" means that it was greater than 20 degrees.

40 [0036] A stain resistance test was conducted by allowing each test sample of Examples 1-1, 1-6 and 1-7 and a glass substrate itself to stand still in the outdoors. After that, the degree of stains on the coated side of the test sample and that on the glass substrate were checked with the naked eyes. With this, the former was found to be considerably less than the latter.

Table 1

	Abrasion Resistance	Acid Resistance	Alkali Resistance	Photo-catalytic Activity	Hydrophilicity	Overall Evaluation
50 Example 1-1	Good	Good	Good	10	Good	Good
Example 1-2	Good	Good	Good	19	Good	Good
Example 1-3	Good	Good	Good	6	Good	Good
Example 1-4	Good	Good	Good	8	Good	Good
55 Example 1-5	Good	Good	Good	9	Good	Good
Example 1-6	Good	Good	Good	32	Good	Good

EP 1 081 108 A1

Table 1 (continued)

	Abrasion Resistance	Acid Resistance	Alkali Resistance	Photo-catalytic Activity	Hydrophilicity	Overall Evaluation
Example 1-7	Good	Good	Good	18	Good	Good
Example 1-8	Good	Good	Good	8	Good	Good
Com. Ex. 1-1	Not good	Good	Not good	48	Good	Not good
Com. Ex. 1-2	Good	Good	Not good	3	Good	Not good
Com. Ex. 1-3	Good	Good	Not good	4	Good	Not good
Com. Ex. 1-4	Not good	Good	Good	20	Good	Not good
Com. Ex. 1-5	Good	Good	Good	2	Good	Not good
Com. Ex. 1-6	Good	Good	Good	3	Good	Not good
Com. Ex. 1-7	Not good	Good	Good	21	Good	Not good
Com. Ex. 1-8	Good	Good	Good	0	Not good	Not good
Com. Ex. 1-9	Good	Good	Good	0	Not good	Not good
Com. Ex. 1-10	Good	Good	Not good	20	Good	Not good

EXAMPLE 1-2

25 [0037] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 40 wt% of ZrO_2 , 20 wt% of SiO_2 and 40 wt% of TiO_2 .

EXAMPLE 1-3

30 [0038] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 30 wt% of ZrO_2 , 40 wt% of SiO_2 and 30 wt% of TiO_2 and that the film had a thickness of 150 nm.

EXAMPLE 1-4

35 [0039] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 50 wt% of ZrO_2 , 20 wt% of SiO_2 and 30 wt% of TiO_2 and that the film had a thickness of 150 nm.

EXAMPLE 1-5

40 [0040] Example 1-1 was repeated except in that a mixture was prepared by dispersing in ethanol MSH2 (trade name) of Mitsubishi Chemical Corp. containing a polymethoxysiloxane as the raw material of SiO_2 and PC-101 (trade name) of Titan Kogyo K.K. as the raw material of the TiO_2 crystals, using a ball mill, and then $Zr(OC_4H_9)_3Cl$ as the raw material of ZrO_2 was added to the mixture, thereby preparing a coating liquid.

EXAMPLE 1-6

45 [0041] Example 1-1 was repeated except in that the coating liquid was applied by dip coating to the surface of a glass plate that is the same as that of Example 1-1, and then the coated glass plate was subjected to a heat treatment by putting it for 40 minutes in a FP41-type muffle furnace of Yamato Kagaku Co. of 450°C, thereby forming a film having a thickness of 80 nm.

EXAMPLE 1-7

50 [0042] Example 1-3 was repeated except in that the coating liquid was applied by dip coating to the surface of a glass plate that is the same as that of Example 1-1, and then the coated glass plate was subjected to a heat treatment by putting it for 40 minutes in a FP41-type muffle furnace of Yamato Kagaku Co. of 450°C.

EXAMPLE 1-8

[0043] Example 1-1 was repeated except in that a coating liquid was prepared by adding aluminum nitrate of Kishida Chemical Co. as the raw material of Al_2O_3 to the coating liquid of Example 1-1 such that the film contained 30 wt% of ZrO_2 , 5 wt% of Al_2O_3 , 25 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-1

[0044] Example 1-1 was repeated except in that a coating liquid was prepared by dispersing only ST-K01 (trade name) of Ishihara Techno Co. in a mixture of ethanol and 1-methoxy-2-propanol each being made of Kishida Chemical Co. such that the film contained 20 wt% of SiO_2 and 80 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-2

[0045] Example 1-1 was repeated except in that a coating liquid was prepared by dispersing only ST-K03 (trade name) of Ishihara Techno Co., in which the weight ratio of TiO_2 to SiO_2 is 5:5, in a mixture of ethanol and 1-methoxy-2-propanol each being made of Kishida Chemical Co. such that the film contained 50 wt% of SiO_2 and 50 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-3

[0046] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 10 wt% of ZrO_2 , 40 wt% of SiO_2 and 50 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-4

[0047] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 30 wt% of ZrO_2 , 20 wt% of SiO_2 and 50 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-5

[0048] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 10 wt% of ZrO_2 , 50 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-6

[0049] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 20 wt% of ZrO_2 , 40 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-7

[0050] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 50 wt% of ZrO_2 , 10 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-8

[0051] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 10 wt% of ZrO_2 , 70 wt% of SiO_2 and 20 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-9

[0052] Example 1-1 was repeated except in that a coating liquid was prepared such that the film contained 30 wt% of ZrO_2 , 50 wt% of SiO_2 and 20 wt% of TiO_2 .

COMPARATIVE EXAMPLE 1-10

[0053] Example 1-1 was repeated except that a coating liquid was prepared by mixing (1) CSG-DI-0600 (trade name) of Chisso Co. as the raw material of SiO_2 , (2) ST-K01 (trade name) of Ishihara Techno Co., in which the weight ratio of TiO_2 to SiO_2 is 8:2, as the raw material of TiO_2 crystals and SiO_2 , and (3) a liquid containing $Ti(OC_3H_7)_4$ which

has been dissolved in ethanol and stabilized with an equimolar amount of acetylacetone, with a mixture of ethanol and 1-methoxy-2-propanol each being made by Kishida Chemical Co. such that the film contained 30 wt% of TiO_2 derived from $\text{Ti}(\text{OC}_3\text{H}_7)_4$, 30 wt% of SiO_2 and 40 wt% of TiO_2 derived from ST-K01.

EXAMPLE 2-1

[0054] A coating liquid was prepared in the same manner as that of Example 1-1. Separately, a float glass plate of soda-lime-silicate glass having dimensions of 1,000mm, 1,800mm and 3.5 mm (thickness) was sufficiently washed with water, detergent and ceria. Then, the coating liquid was applied to the glass plate by dip coating, followed by a preliminary baking at 600°C for 5 minutes. Then, the coated glass plate was cut into a predetermined shape, then subjected to several preliminary treatments, such as seaming (chamfering), the application of a black paint to the periphery, a heating wire printing and the like, and then subjected to a main baking at 650°C, thereby obtaining an article of the invention, that is, an automotive rear window glass pane with a photocatalytic film having a thickness of 120 nm and a composition of 30 wt% of ZrO_2 , 30 wt% of SiO_2 and 40 wt% of TiO_2 , based on the weight of this film. During the main baking, the glass plate was bent into a curved shape.

[0055] The obtained article (test sample) was subjected to the following evaluation tests. The results of the tests are shown in Table 2.

[0056] An abrasion resistance test was conducted in the same manner as that of Example 1-1. In Table 2, "good" means that the haze value before the test H_0 is 1% or less, and ΔH is 2% or less where $\Delta H = H_{\text{max}} - H_0$ in which H_{max} is the maximum haze value during 1,000 cycles. In Table 2, "not good" means that H_0 is greater than 1% or ΔH is greater than 2%.

[0057] An acid resistance test was conducted in the same manner as that of Example 1-1, except in that 1N hydrochloric acid was replaced with 1 wt% sulfuric acid. The evaluation was conducted in the same manner as that of Example 1-1.

[0058] An alkali resistance test was conducted in the same manner as that of Example 1-1, except in that 1N sodium hydroxide aqueous solution was replaced with 1wt% sodium hydroxide aqueous solution. The evaluation was conducted in the same manner as that of Example 1-1.

[0059] A photocatalytic activity test and its evaluation were conducted in the same manners as those of Example 1-1.

[0060] A hydrophilicity test and its evaluation were conducted in the same manners as those of Example 1-1.

[0061] A stain resistance test was conducted by allowing the test sample of Example 2-1 and an automotive rear window glass having no photocatalytic film thereon to stand still in the outdoors. After that, the degree of stains on the coated side of the test sample and that on the automotive rear window glass were checked with the naked eyes. With this, the former was found to be considerably less than the latter.

Table 2

	Abrasion Resistance	Acid Resistance	Alkali Resistance	Photocatalytic Activity	Hydrophilicity	Overall Evaluation
Example 2-1	Good	Good	Good	14	Good	Good
Example 2-2	Good	Good	Good	19	Good	Good
Example 2-3	Good	Good	Good	6	Good	Good
Example 2-4	Good	Good	Good	8	Good	Good
Example 2-5	Good	Good	Good	9	Good	Good
Example 2-6	Good	Good	Good	8	Good	Good
Com. Ex. 2-1	Not good	Good	Not good	48	Good	Not good
Com. Ex. 2-2	Good	Good	Not good	3	Good	Not good
Com. Ex. 2-3	Good	Good	Not good	4	Good	Not good
Com. Ex. 2-4	Not good	Good	Good	20	Good	Not good
Com. Ex. 2-5	Good	Good	Good	2	Good	Not good
Com. Ex. 2-6	Good	Good	Good	3	Good	Not good

Table 2 (continued)

	Abrasion Resistance	Acid Resistance	Alkali Resistance	Photocatalytic Activity	Hydrophilicity	Overall Evaluation
5 Com. Ex. 2-7	Not good	Good	Good	21	Good	Not good
Com. Ex. 2-8	Good	Good	Good	0	Not good	Not good
Com. Ex. 2-9	Good	Good	Good	0	Not good	Not good
10 Com. Ex. 2-10	Good	Good	Not good	20	Good	Not good

EXAMPLE 2-2

15 [0062] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 40 wt% of ZrO_2 , 20 wt% of SiO_2 and 40 wt% of TiO_2 .

EXAMPLE 2-3

20 [0063] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 30 wt% of ZrO_2 , 40 wt% of SiO_2 and 30 wt% of TiO_2 and that the film had a thickness of 150 nm.

EXAMPLE 2-4

25 [0064] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 50 wt% of ZrO_2 , 20 wt% of SiO_2 and 30 wt% of TiO_2 and that the film had a thickness of 150 nm.

EXAMPLE 2-5

30 [0065] Example 2-1 was repeated except in that a mixture was prepared by dispersing in ethanol MSH2 (trade name) of Mitsubishi Chemical Corp. as the raw material of SiO_2 and PC-101 (trade name) of Titan Kogyo K.K. as the raw material of the TiO_2 crystals, using a ball mill, and then $Zr(OC_4H_9)_3Cl$ as the raw material of ZrO_2 was added to the mixture, thereby preparing a coating liquid.

EXAMPLE 2-6

35 [0066] Example 2-1 was repeated except in that a coating liquid was prepared by adding aluminum nitrate of Kishida Chemical Co. as the raw material of Al_2O_3 to the coating liquid of Example 1-1 such that the film contained 30 wt% of ZrO_2 , 5 wt% of Al_2O_3 , 25 wt% of SiO_2 and 40 wt% of TiO_2 .

40 COMPARATIVE EXAMPLE 2-1

[0067] Example 2-1 was repeated except in that a coating liquid was prepared by dispersing only ST-K01 (trade name) of Ishihara Techno Co. in a mixture of ethanol and 1-methoxy-2-propanol each being made of Kishida Chemical Co. such that the film contained 20 wt% of SiO_2 and 80 wt% of TiO_2 .

45

COMPARATIVE EXAMPLE 2-2

[0068] Example 2-1 was repeated except in that a coating liquid was prepared by dispersing only ST-K03 (trade name) of Ishihara Techno Co., in which the weight ratio of TiO_2 to SiO_2 is 5:5, in a mixture of ethanol and 1-methoxy-2-propanol each being made of Kishida Chemical Co. such that the film contained 50 wt% of SiO_2 and 50 wt% of TiO_2 .

50

COMPARATIVE EXAMPLE 2-3

55 [0069] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 10 wt% of ZrO_2 , 40 wt% of SiO_2 and 50 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-4

[0070] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 30 wt% of ZrO_2 , 20 wt% of SiO_2 and 50 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-5

[0071] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 10 wt% of ZrO_2 , 50 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-6

[0072] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 20 wt% of ZrO_2 , 40 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-7

[0073] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 50 wt% of ZrO_2 , 10 wt% of SiO_2 and 40 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-8

[0074] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 10 wt% of ZrO_2 , 70 wt% of SiO_2 and 20 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-9

[0075] Example 2-1 was repeated except in that a coating liquid was prepared such that the film contained 30 wt% of ZrO_2 , 50 wt% of SiO_2 and 20 wt% of TiO_2 .

COMPARATIVE EXAMPLE 2-10

[0076] Example 2-1 was repeated except that a coating liquid was prepared by mixing (1) CSG-DI-0600 (trade name) of Chisso Co. as the raw material of SiO_2 , (2) ST-K01 (trade name) of Ishihara Techno Co., in which the weight ratio of TiO_2 to SiO_2 is 8:2, as the raw material of TiO_2 crystals and SiO_2 , and (3) a liquid containing $Ti(OC_3H_7)_4$ which has been dissolved in ethanol and stabilized with an equimolar amount of acetylacetone, with a mixture of ethanol and 1-methoxy-2-propanol each being made by Kishida Chemical Co. such that the film contained 30 wt% of TiO_2 derived from $Ti(OC_3H_7)_4$, 30 wt% of SiO_2 and 40 wt% of TiO_2 derived from ST-K01.

[0077] The entire disclosure of each of Japanese Patent Application Nos. 11-248155 filed on September 2, 1999 and 11-367121 filed on December 24, 1999, including specification, claims and summary, is incorporated herein by reference in its entirety.

Claims

1. An article comprising:

a substrate; and
a photocatalytic film covering said substrate, said film comprising (1) a film-forming component comprising ZrO_2 and SiO_2 and (2) TiO_2 crystals dispersed in said film-forming component, said ZrO_2 , said SiO_2 and said TiO_2 crystals respectively being in amounts of 25-60 wt%, 15-50 wt% and 25-45 wt%, based on a weight of said film.

2. An article according to claim 1, wherein a total amount of said ZrO_2 , said SiO_2 and said TiO_2 crystals is at least 90 wt%, based on the weight of said film.

3. An article according to claim 1, wherein said film has a thickness of from 50 to 300 nm.

4. An article according to claim 1, wherein said ZrO_2 is in an amount of 25-45 wt%, based on the weight of said film.

5. An article according to claim 1, wherein said SiO_2 is in an amount of 25-45 wt%, based on the weight of said film.
6. An article according to claim 1, wherein said TiO_2 crystals have an average grain size of not greater than 30 nm.
- 5 7. An article according to claim 6, wherein said average grain size is not greater than 20 nm.
8. An article according to claim 1, wherein said substrate is a curved glass plate.
9. An article according to claim 1, wherein said article is an automotive window pane having said photocatalytic film
10 on an outer side thereof.
10. An article according to claim 1, wherein said film further comprises at least one member selected from the group
consisting of amorphous TiO_2 , colloidal silica, Al_2O_3 , B_2O_3 , SnO_2 , Cr ion, V ion, and carbon.
- 15 11. An article according to claim 1, wherein said article is produced by a process comprising:

providing a coating liquid comprising a first raw material of said ZrO_2 , a second raw material of said SiO_2 , and
a third raw material of said TiO_2 crystals;
applying said coating liquid to a surface of said substrate, thereby forming thereon a precursory film ; and
20 subjecting said precursory film to a heat treatment at a temperature of 300-700°C, thereby turning said precursory
film into said photocatalytic film.
12. An article according to claim 11, wherein said third raw material is TiO_2 crystal grains having an average grain size
of not greater than 20 nm.
- 25 13. An article according to claim 11, wherein said first raw material is zirconium chloride or zirconium nitrate.
14. An article according to claim 1, wherein said article is produced by a process comprising:

30 providing a coating liquid comprising a first raw material of said ZrO_2 , a second raw material of said SiO_2 , and
a third raw material of said TiO_2 crystals;
applying said coating liquid to a surface of said substrate, thereby forming thereon a precursory film ; and
subjecting said precursory film to a first baking at a temperature of 560-700°C, thereby turning said precursory
35 film into said photocatalytic film.
15. An article according to claim 14, wherein said substrate is a glass plate, and wherein said glass plate is bent into a
curved glass plate when said precursory film and said glass plate are subjected to said first baking.
16. An article according to claim 14, wherein said substrate is a glass plate, and wherein said glass plate is bent into a
40 curved glass plate after said first baking.
17. An article according to claim 14, wherein, prior to said first baking, said precursory film is subjected to a second
baking at a temperature of 300-620°C.
- 45 18. An article according to claim 14, wherein said third raw material is TiO_2 crystal grains having an average grain size
of not greater than 30 nm.
19. An article according to claim 14, wherein said first raw material is zirconium chloride or zirconium nitrate.
- 50 20. A process for producing an article, said article comprising:

a substrate; and
a photocatalytic film covering said substrate, said film comprising (1) a film-forming component comprising
55 ZrO_2 and SiO_2 and (2) TiO_2 crystals dispersed in said film-forming component, said ZrO_2 , said SiO_2 and said
 TiO_2 crystals respectively being in amounts of 25-60 wt%, 15-50 wt% and 25-45 wt%, based on a weight of
said film,
said process comprising:
providing a coating liquid comprising a first raw material of said ZrO_2 , a second raw material of said SiO_2 , and

a third raw material of said TiO_2 crystals;
 applying said coating liquid to a surface of said substrate, thereby forming thereon a precursory film; and
 subjecting said precursory film to a heat treatment at a temperature of 300-700°C, thereby turning said precursory film into said photocatalytic film.

21. A process according to claim 20, wherein said third raw material is TiO_2 crystal grains having an average grain size of not greater than 20 nm.

22. A process according to claim 20, wherein said first raw material is zirconium chloride or zirconium nitrate.

23. A process for producing an article, said article comprising:

a substrate; and

a photocatalytic film covering said substrate, said film comprising (1) a film-forming component comprising ZrO_2 and SiO_2 and (2) TiO_2 crystals dispersed in said film-forming component, said ZrO_2 , said SiO_2 and said TiO_2 crystals respectively being in amounts of 25-60 wt%, 15-50 wt% and 25-45 wt%, based on a weight of said film,

said process comprising:

providing a coating liquid comprising a first raw material of said ZrO_2 , a second raw material of said SiO_2 , and a third raw material of said TiO_2 crystals;

applying said coating liquid to a surface of said substrate, thereby forming thereon a precursory film; and
 subjecting said precursory film to a first baking at a temperature of 560-700°C, thereby turning said precursory film into said photocatalytic film.

24. A process according to claim 23, wherein said substrate is a glass plate, and wherein said glass plate is bent into a curved glass plate when said precursory film and said glass plate are subjected to said first baking.

25. A process according to claim 23, wherein said substrate is a glass plate, and wherein said glass plate is bent into a curved glass plate after said first baking.

26. A process according to claim 23, wherein, prior to said first baking, said precursory film is subjected to a second baking at a temperature of 300-620°C.

27. A process according to claim 23, wherein said third raw material is TiO_2 crystal grains having an average grain size of not greater than 30 nm.

28. A process according to claim 23, wherein said first raw material is zirconium chloride or zirconium nitrate.



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 00 11 8871

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	EP 0 911 300 A (CENTRAL GLASS CO LTD) 28 April 1999 (1999-04-28) * claims; example 3 *	1-28	C03C17/23 C03C17/245 C03C17/25
X	FR 2 738 812 A (SAINT GOBAIN VITRAGE) 21 March 1997 (1997-03-21) * claims 1-11 *	1-3	
D,X	DATABASE WPI Section Ch, Week 199843 Derwent Publications Ltd., London, GB; Class J04, AN 1998-499640 XP002155216 & JP 10 216528 A (SHARP KK), 18 August 1998 (1998-08-18) * abstract *	1-3	
D,X	DATABASE WPI Section Ch, Week 199810 Derwent Publications Ltd., London, GB; Class L01, AN 1998-105013 XP002155217 & JP 09 328336 A (ASAHI GLASS CO LTD), 22 December 1997 (1997-12-22) * abstract *	1-28	TECHNICAL FIELDS SEARCHED (Int.Cl.7) C03C
A	FR 2 738 836 A (RHONE POULENC CHIMIE) 21 March 1997 (1997-03-21) * claims *	1-28	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 12 December 2000	Examiner Reedijk, A
CATEGORY OF CITED DOCUMENTS		T: theory or principle underlying the invention E: earlier patent document, but published on, or after the filing date D: document cited in the application L: document cited for other reasons A: technological background O: non-written disclosure P: intermediate document &: member of the same patent family, corresponding document	
X: particularly relevant if taken alone Y: particularly relevant if combined with another document of the same category A: technological background O: non-written disclosure P: intermediate document			

EPO FORM 1603/03 02 (P4/C31)



EP 1 081 108 A1

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 11 8871

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

12-12-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0911300 A	28-04-1999	JP 11130468 A	18-05-1999
		JP 11180733 A	06-07-1999
		US 6055085 A	25-04-2000
FR 2738812 A	21-03-1997	AU 6992796 A	01-04-1997
		BR 9610289 A	21-12-1999
		CZ 9800756 A	12-08-1998
		EP 0850203 A	01-07-1998
		WO 9710185 A	20-03-1997
		JP 11512336 T	26-10-1999
		PL 325526 A	03-08-1998
		US 6037289 A	14-03-2000
JP 10216528 A	18-08-1998	NONE	
JP 9328336 A	22-12-1997	NONE	
FR 2738836 A	21-03-1997	AU 6992796 A	01-04-1997
		BR 9610289 A	21-12-1999
		CZ 9800756 A	12-08-1998
		EP 0850203 A	01-07-1998
		WO 9710185 A	20-03-1997
		JP 11512336 T	26-10-1999
		PL 325526 A	03-08-1998
		US 6037289 A	14-03-2000

EPO FORM P0468

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82